

ABSTRACT

A low toxicity, stable oxazaphosphorine containing compositions with mesna for parenteral administration has been described. The process essentially requires addition of an oxazaphosphorine antineoplastic to the aqueous solution of an etherified β -cyclodextrin followed by addition of mesna as such or as an aqueous solution containing optionally, an etherified β -cyclodextrin. Preferably, the oxazaphosphorine antineoplastic is Ifosfamide and the etherified β -cyclodextrin is 2-hydroxypropyl- β -cyclodextrin.

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